S

	Application Number		10601171	
	Filing Date		2003-06-23	
NFORMATION DISCLOSURE TATEMENT BY APPLICANT Not for submission under 37 CFR 1.99)	First Named Inventor Geral		ld Walter FISCHER	
	Art Unit		1645	
	Examiner Name	N. Archie		
	Attorney Docket Numb	or	SYNL003CNRCE	

U.S.PATENTS					Remove		
Examiner Cite Patent Number Kind Code1 Issue Date Name of Patentee or Applica of cited Document				Name of Patentee or Applicant of cited Document	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		
	1						
If you wisl	h to add	additional U.S. Pater	t citatio	n information pl	ease click the Add button.	Add	
			U.S.P	ATENT APPLI	CATION PUBLICATIONS	Remove	
Examiner Initial* Cite No Publication Number Code Date No Publication Name of Patentee or Applicant Passages or Releving Technology (Code Date No Name of Patentee or Applicant Passages or Releving Name of Cited Document Passages or Releving Name of Patentee or Applicant Passages or Patentee or Passages or Patentee or							
	1						
If you wisl	h to add	additional U.S. Publis	hed Ap	plication citation	n information please click the Ad	d button. Add	
					THE PARTITION	Demoire	

FOREIGN PATENT DOCUMENTS						ENIS	TOHIOTO		
Examine Initial*	Cite No	Foreign Document Number ³	Country Code ² i	Kind Code4	Publication	Name of Patentee or Applicant of cited Document	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		
	1	93/06213	wo	A1	1993-04-01	Medical Research Council et al.			

If you wish to add additional Foreign Patent Document citation information please click the Add button Add

NON-PATENT LITERATURE DOCUMENTS Remove

NON-PATENT LITERATURE DOCUMENTS Remove

Non-PATENT LITERATURE DOCUMENTS Remove

initials* No | (block marpaine, journal serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), pages(s), volume

1	European Office Action for Application No. 10182580.0, dated July 4, 2011	

If you wish to add additional non-patent literature document citation information please click the Add button Add

EXAMINER SIGNATURE

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

Date Considered

See Kinz Code of UBPTO Please Documents at invest UBSTO_CGLY or MEDF 90.04. If Sets or files that issued the document, by the located are code (NIPO) shaded at 73... If you please parted for committed the or files of the please that process the sets in them or the please to the please that process the sets in them or the please to the please that process the sets in them or the please to make the please that th

Examiner Signature

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number Filing Date First Named Inventor Gerald Art Unit			10601171		
			2003-06-23		
		Geral	d Walter FISCHER		
			1645		
	Examiner Name	N. Arc	thie		
Attorney Docket Number		er	SYNI-003CNRCE		

CERTIFICATION STATEMENT

Please see 37	CFR 1.9	7 and 1.98 to	make the appro	priate selection(s	1
---------------	---------	---------------	----------------	--------------------	---

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. Sea 37 CFF 1.37(e)(1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any involved designated in 37 CFR 1.56(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.97(e)(c).

- See attached certification statement.
- The fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- A certification statement is not submitted herewith

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Amy E. Mandragouras, Esq./	Date (YYYY-MM-DD)	2011-08-25		
Name/Print	Amy E. Mandragouras, Esq.	Registration Number	36207		

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file fand by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C. 12.0 and 3T CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete his form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenary Cfiler. U.S. Operatment of Commence, P. 0. Box 1450, Alexandria, V.S. 2313-1450. D. ONT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.S. 2313-1450.

Privacy Act Statement

The Privacy Act of 1974 (P. L. 95.79) requires that you be given certain information in connection with your submission of the stackhold from reliable to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. (2)(2)(2) familishing of the information solicided is columbra; and (3) the principal purpuse for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or cosmitine your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or cosmitine your submission related to a patient agricultant or patient. If you do not furnish the requested requirement of the patient of the pati

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
- A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
- A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the sublect matter of the record.
- A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, ourspant to 5 U.S.C. S52a(m).
- A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
- A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designee, during an inspection of records conducted by GSAs a part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2504 and 2506. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- 8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application presume to 58 U.S. C. 12(p) or issuance of a patent pursuant to 58 U.S. C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 11, 4, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by whiter a published application, an application open to public inscredience or an issued patent.
- A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.